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SYNTHESIS OF NANOSTRUCTURED MnO_x FOR LOW-TEMPERATURE NO_x SCR / Andreoli, Silvia; Deorsola, FABIO ALESSANDRO; Pirone, Raffaele. - (2013). (Intervento presentato al convegno XVII National Congress of Catalysis GIC 2013 tenutosi a Riccione (IT) nel 15-18 settembre 2013).

Availability:

This version is available at: 11583/2634851 since: 2016-02-23T14:59:22Z

Publisher:

Published

DOI:

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SYNTHESIS OF NANOSTRUCTURED MnO_x FOR LOW-TEMPERATURE NO_x SCR

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Nitrogen oxides (NO , N_2O and NO_2) are by-produced and emitted in high-temperature combustion of stationary and mobile sources and constitute well-known atmospheric pollutants that can contribute to the ozone depletion, acids rains, photochemical smog and greenhouse effects.

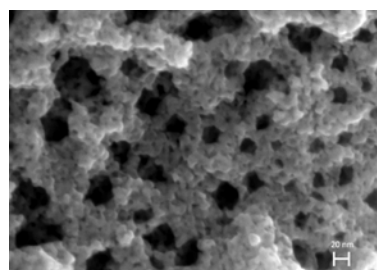
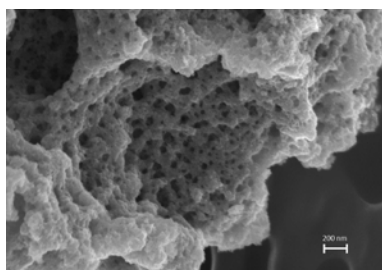
In large stationary applications (power plants, chemical industry, etc.), the issue of NO_x removal is accomplished through the post-treatment of the exhaust gases by means of the selective catalytic reduction (SCR) method. Such a technology should be now revised to perform the abatement on nitrogen oxides in mobile sources too, since especially for diesel engines no solution seems to be viable with non-specific reductants (CO , HCs), with the consequent necessity to develop active catalysts for the reduction of NO_x with NH_3 in a significantly lower temperature range.

In the present work, the properties of manganese oxide (MnO_x) based catalysts have been deeply investigated, because manganese arouses a great research interest in low temperature SCR, but the exact nature of active sites seems not to have been clarified, neither the optimal oxidation state of the metal for the reaction has been assessed.

The synthesis of the MnO_x catalysts was carried out through the Solution Combustion Synthesis (SCS) method, that was demonstrated to be effective in the achievement of nanostructured catalysts with high specific surface area [1,2]. The method is based on an exothermic oxidation-reduction reaction between suitable amounts of a salt of the desired metal oxide and an organic combustible, in this case manganese nitrate and glycine.

A series of characterization analyses have been carried out over different MnO_x samples, prepared with the aim of having pure oxides phases or at least a variable distribution of the oxidation state of Mn. The structural characterization by X-ray diffraction (XRD) showed, for all samples, peaks belonging to the Mn_2O_3 phase, that resulted narrower and sharper by increasing the synthesis temperature, meaning an increased crystallinity of the catalysts.

The morphological characterization, performed by FESEM observations, is represented in the Figure below. The catalysts showed the presence of primary nanoparticles, with size of about 20 nm (see Figure on the right), disposed into large agglomerates characterized by a significant amount of porosity, with pore dimension in the range between 20 and 100 nm. The value of specific surface area of the samples, measured by the BET method, was $64.5 \text{ m}^2/\text{g}$.



Moreover, TPR analyses have been carried out over all samples and put in connection with the results of catalytic activity tests for the SCR reaction with ammonia.

References.

- [1] Mescia, D., Caroca, J.C., Russo, N., Labhsetwar, N., Fino, D., Saracco, G., Specchia, V. (2008): *Catal. Today*, **137**, 300-305;
- [2] Russo, N., Fino, D., Saracco, G., Specchia, V. (2006): *Catal. Today*, **117**, 214–219.